

University of Groningen

Decision aids that facilitate elements of shared decision making in chronic illnesses

Wieringa, Thomas H; Rodriguez-Gutierrez, Rene; Spencer-Bonilla, Gabriela; de Wit, Maartje; Ponce, Oscar J; Sanchez-Herrera, Manuel F; Espinoza, Nataly R; Zisman-Ilani, Yaara; Kunneman, Marleen; Schoonmade, Linda J

Published in:
Systematic Reviews

DOI:
[10.1186/s13643-019-1034-4](https://doi.org/10.1186/s13643-019-1034-4)

IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.

Document Version
Publisher's PDF, also known as Version of record

Publication date:
2019

[Link to publication in University of Groningen/UMCG research database](#)

Citation for published version (APA):

Wieringa, T. H., Rodriguez-Gutierrez, R., Spencer-Bonilla, G., de Wit, M., Ponce, O. J., Sanchez-Herrera, M. F., Espinoza, N. R., Zisman-Ilani, Y., Kunneman, M., Schoonmade, L. J., Montori, V. M., & Snoek, F. J. (2019). Decision aids that facilitate elements of shared decision making in chronic illnesses: a systematic review. *Systematic Reviews*, 8(1), [121]. <https://doi.org/10.1186/s13643-019-1034-4>

Copyright

Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

The publication may also be distributed here under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license. More information can be found on the University of Groningen website: <https://www.rug.nl/library/open-access/self-archiving-pure/taverne-amendment>.

Take-down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.


Downloaded from the University of Groningen/UMCG research database (Pure): <http://www.rug.nl/research/portal>. For technical reasons the number of authors shown on this cover page is limited to 10 maximum.

RESEARCH

Open Access



Decision aids that facilitate elements of shared decision making in chronic illnesses: a systematic review

Thomas H. Wieringa^{1*} , Rene Rodriguez-Gutierrez^{2,3,4}, Gabriela Spencer-Bonilla^{2,5}, Maartje de Wit¹, Oscar J. Ponce², Manuel F. Sanchez-Herrera², Nataly R. Espinoza², Yaara Zisman-Ilani⁶, Marleen Kunneman^{2,7}, Linda J. Schoonmade⁸, Victor M. Montori² and Frank J. Snoek¹

Abstract

Background: Shared decision making (SDM) is a patient-centered approach in which clinicians and patients work together to find and choose the best course of action for each patient's particular situation. Six SDM key elements can be identified: situation diagnosis, choice awareness, option clarification, discussion of harms and benefits, deliberation of patient preferences, and making the decision. The International Patient Decision Aid Standards (IPDAS) require that a decision aid (DA) support these key elements. Yet, the extent to which DAs support these six key SDM elements and how this relates to their impact remain unknown.

Methods: We searched bibliographic databases (from inception until November 2017), reference lists of included studies, trial registries, and experts for randomized controlled trials of DAs in patients with cardiovascular, or chronic respiratory conditions or diabetes. Reviewers worked in duplicate and independently selected studies for inclusion, extracted trial, and DA characteristics, and evaluated the quality of each trial.

Results: DAs most commonly clarified options (20 of 20; 100%) and discussed their harms and benefits (18 of 20; 90%; unclear in two DAs); all six elements were clearly supported in 4 DAs (20%). We found no association between the presence of these elements and SDM outcomes.

Conclusions: DAs for selected chronic conditions are mostly designed to transfer information about options and their harms and benefits. The extent to which their support of SDM key elements relates to their impact on SDM outcomes could not be ascertained.

Systematic review registration: PROSPERO registration number: [CRD42016050320](https://www.crd42016050320).

Keywords: Chronic illnesses, Decision aids, Shared decision making

Background

Shared decision making (SDM) is a patient-centered approach in which clinicians and patients work together to find and choose (by taking into account the best available evidence, as well as the patients' problems, values, preferences, and contexts) the best course of action for

each patient's particular situation [1], an approach that is pertinent to the care of patients with chronic conditions [2]. Decisions in the context of self-management of chronic conditions differ from one-time decisions, as in the former decisions can often be reconsidered [2]. Six key elements of SDM can be identified from the literature: situation diagnosis, choice awareness, option clarification, discussion of harms and benefits, patient preferences deliberation, and making the decision [1–4]. As noted by Stiggelbout and others [5, 6], SDM promotes actions that are needed, wanted, and more likely to be implemented. A shared understanding and treatment focused on patients' health and life goals, as well

* Correspondence: t.wieringa@vumc.nl

¹Department of Medical Psychology, Amsterdam Public Health Research Institute, Amsterdam UMC, Vrije Universiteit Amsterdam, De Boelelaan 1117, Amsterdam, the Netherlands

Full list of author information is available at the end of the article



© The Author(s). 2019 **Open Access** This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (<http://creativecommons.org/licenses/by/4.0/>), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The Creative Commons Public Domain Dedication waiver (<http://creativecommons.org/publicdomain/zero/1.0/>) applies to the data made available in this article, unless otherwise stated.

as a stronger clinician-patient relationship, may also be facilitated by SDM [7, 8].

An SDM interaction starts with a diagnostic conversation (situation diagnosis) [1]. This opening first focuses on understanding the patient's situation and establishing the aspects that require action [1, 4]. When multiple reasonable options are available, then the clinician should indicate this and highlight the importance of the patient's preferences in deciding on the course of action (choice awareness) [3]. Subsequently, the patient and clinician deliberate about the way each option fits and accommodates within each patient's situation (option clarification, discussion of harms and benefits, and patient preferences deliberation). Finally, a decision is made by the clinician and patient (making the decision) [2, 4].

SDM can be facilitated by decision aids (DAs) that have been developed for use by clinicians and patients, either during or in preparation for the clinical encounter [9–11]. DAs can help patients choose an option that is congruent with their values, reduce the proportion of patients remaining undecided and/or who play a passive role in the decision-making process, and improve patient knowledge, decisional conflict, and patient-clinician communication [11–15]. The International Patient Decision Aid Standards (IPDAS) Collaboration aims to enhance the quality and effectiveness of DAs by establishing an evidence-informed framework for improving their content, development, implementation, and evaluation [16]. The IPDAS Collaboration defines a DA as “a tool designed to help people participate in decision making about health care options” [9], and developed a minimal set of standards for qualifying a tool as a DA [17]. According to this minimal set, all SDM key elements, except making the decision, should be handled by a tool in order to regard it as a DA [17]. Despite this minimal set of qualifying criteria, investigators have found that fostering choice awareness through the use of a DA was not a prerequisite for fostering choice awareness per se during the encounter [18]. Therefore, it is unclear whether tools should support all qualifying IPDAS criteria for these tools to support SDM. Therefore, we define a DA in the current review as “any tool designed to support SDM.”

To the best of our knowledge, there is no empirical data to tell us which of the six key elements are supported by DAs and whether there is an association between support for these key elements and SDM outcomes. We hypothesize that DAs that cover multiple elements of SDM are more likely to have positive effects on SDM outcomes, as well as on patient-reported outcomes (PROs). With regard to surrogate and clinical outcomes, there is no reason to expect a consistent response. A previous systematic review of the effects of DAs found that more detailed DAs better improve

knowledge and reduce some aspects of decisional conflict compared to simple DAs, and concluded that more research is needed to evaluate the level of detail needed in DAs [19]. The current review aims to meet this need by studying the SDM elements incorporated in DAs and their effect on SDM outcomes.

This review aims to (1) describe the SDM elements present in DAs for patients with common chronic conditions (cardiovascular, chronic respiratory diseases or diabetes) tested in randomized controlled trials (RCTs), and (2) determine an association between the key elements present and the effects of these DAs compared to usual care or active controls on SDM outcomes (e.g., conversation duration, patient participation, knowledge, and decisional conflict).

Methods

The protocol of this systematic review was previously published [20] and registered in the International Prospective Register of Systematic Reviews (PROSPERO registration number: CRD42016050320; <http://www.crd.york.ac.uk/PROSPERO>). The review is reported according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines [21]. Additional file 1 provides the PRISMA checklist.

Study eligibility

We searched RCTs comparing the use of DAs (any tool designed to support SDM) to usual care or active controls (except other DAs) in adults with cardiovascular disease, chronic respiratory disease, or diabetes and measuring their impact on SDM and health outcomes (patient-reported, surrogate, and clinical outcomes). As described in detail previously [20], we selected chronic conditions that are most prevalent according to the World Health Organization [22–24], most likely to require self-management, and for which decisions may be revisited. We included all pertinent publications of an eligible study. There were no exclusions based on language or year of publication.

Information sources and search strategy

To identify all relevant publications, we performed systematic searches, in collaboration with a medical librarian (LJS) in the bibliographic databases PubMed, Embase.com, Web of Science, CINAHL (through EBSCO), PsycINFO (through EBSCO), and the Cochrane Library from inception to November 7th, 2017. Search terms included MeSH in PubMed, Emtree in Embase.com, CINAHL headings in CINAHL, indexed terms from the Thesaurus in PsycINFO, and free text terms. We used free text only in the Cochrane Library and Web of Science. Search terms compressing “shared decision making” were used in combination with “cardiovascular

diseases” OR “chronic respiratory diseases” OR “diabetes.” Search results were limited to RCTs. Duplicate articles were excluded. All languages were accepted. The full search strategies for all databases can be found in Additional file 2. In early 2017, THW contacted by e-mail and queried SDM experts participating in the Facebook group “Shared@ Shared Decision Making Network,” and in the LinkedIn groups “Platform SDM GB” and “Shared Decision Making in Netherlands” for additional eligible studies. THW also reviewed trial registries including <http://isrctn.com>, <http://narcis.nl>, <http://trialregister.nl>, and <http://www.clinicaltrials.gov>. MFSH reviewed the reference lists from included studies.

Study selection process

After deduplication, pairs of reviewers (two hired persons, GS-B, RRG, and THW), working independently and in duplicate, assessed each abstract for eligibility. Studies considered potentially eligible by at least one reviewer were included for the full-text phase. THW and RRG reviewed selected full-text articles independently and in duplicate. Disagreements were resolved by a third reviewer (GS-B or OJP).

Data collection process

Data about study and DA characteristics, study quality, and outcomes were extracted by pairs of reviewers working in duplicate (two hired persons, RRG, MFSH, YZI, and THW) with conflict resolved by a third reviewer (GS-B, NRE, YZI, and RRG; YZI and RRG resolved conflicts of parts for which they did not collect data). We used the definitions in Table 1 to determine which key SDM components were present. Sets of three articles were used to train and calibrate reviewers through

extraction and discussion of results among reviewers. Outcomes collected were those most proximate to the encounter of interest.

Risk of bias in individual studies

OJP and THW independently assessed the risk of bias on outcome level at all domains of the Cochrane Collaboration’s tool for RCTs [25, 26], with disagreement resolved by consensus. Because blinding of patients and clinicians to the use of conversation aids is not possible, we ignored the two blinding factors. Otherwise, when one or more of the five other domains was regarded as being at high risk of bias, then the summary assessment of the risk of bias was “high.” If one or more domain was “unclear” and all others were “low risk,” then we summarized the risk of bias as “unclear.” If all domains were “low risk,” then the summary assessment of the risk of bias was “low.”

Outcomes and data synthesis

Data on both SDM (e.g., conversation duration, patient participation, knowledge, and decisional conflict) and health outcomes (patient-reported, surrogate, and clinical outcomes) were collected. Standardized mean differences (SMDs) together with their 95% confidence intervals (95%-CIs) were calculated for continuous outcomes using Review Manager 5.3 [27]. Odds ratios (ORs) together with their 95% CIs were directly extracted from the reports. If the mean difference and/or its standard error (SE) and 95% CI were not presented in the article, then the SMD together with its 95% CI were calculated by entering the mean score/value per arm together with their standard deviations (SDs). If the 95% CI for an OR was not presented, then numbers for every cell in the 2 × 2 table were inserted into Review Manager 5.3 to be calculated. The SMD could not be calculated when only interquartile ranges were reported. We also summarize the data narratively according to our protocol [20].

Missing data and author contact

All corresponding authors (or other authors if no response after approximately 6 weeks) of included studies were contacted through e-mail and, if no response, again approximately 4 weeks later (although originally planned, we did not contact authors by phone) to request missing data or clarifications. If the authors did not respond or could not provide a missing standard deviation needed to calculate the SMD, then the SD of the most comparable study with the same outcome and measurement instrument was imputed.

Table 1 Definitions for the key elements of SDM in decision aids (DAs)

Key element of SDM	Definitions for this study [4, 18]
Situation diagnosis	The DA explicitly describes the patient’s problem.
Choice awareness	The DA explicitly acknowledges that the patient’s situation is mutable, that there is more than one sensible way to address or change this situation, and that patient input matters in deciding how to proceed.
Option clarification	The DA explicitly lists and describes the options available.
Harms and benefits discussion	The DA explicitly explains the harms and benefits of the available options.
Patient preferences deliberation	The DA explicitly elicits the patient’s preferences or explicitly motivates the parties to discuss them.
Making the decision	The DA explicitly elicits the patient’s wish to make or defer a decision, asks for the patient’s choice, or describes the patient’s choice.

Results

Figure 1 describes the flow of the study selection. Chance-adjusted inter-reviewer agreement (k) for eligibility was only fair ($k = 0.3\text{--}0.4$) [28]. We found 24 articles reporting on 23 RCTs of 20 DAs (10 DAs for cardiovascular disease, 2 DAs for respiratory diseases, and 8 DAs for diabetes). The effectiveness of Statin Choice was studied in three RCTs described in four articles meeting our criteria and The Diabetes Medication Choice Decision Aid was studied in two RCTs described in two separate articles. Other DAs were studied in one RCT described in one article. Additional file 3 presents the risk of bias assessment on the outcome level per study. Besides the study of Gagné et al. [29], all studies have an unclear or high risk of bias for all outcomes assessed in this review.

Table 2 shows the SDM elements supported per DA. The elements were described as “unclear” if the DAs were neither described clearly nor available for our inspection, and/or if reviewers were uncertain whether the regarding element was included in the DA. The option clarification element (included in 20 of 20 DAs; 100%) and the harms and benefits discussion (included in 18 of 20 DAs; 90%; unclear in two DAs) are the elements most commonly clearly included in the DAs. The other elements are less common and more uncertainty is present whether these elements are included, especially with regard to choice awareness (uncertain in 14 out of 20 DAs; 70%). All elements were clearly supported in four DAs (20%). Table 2 also shows the DA effects on SDM outcomes. We could not glean any association between SDM elements present in the DAs and SDM outcomes. Additional file 4 reports details of the DAs included here

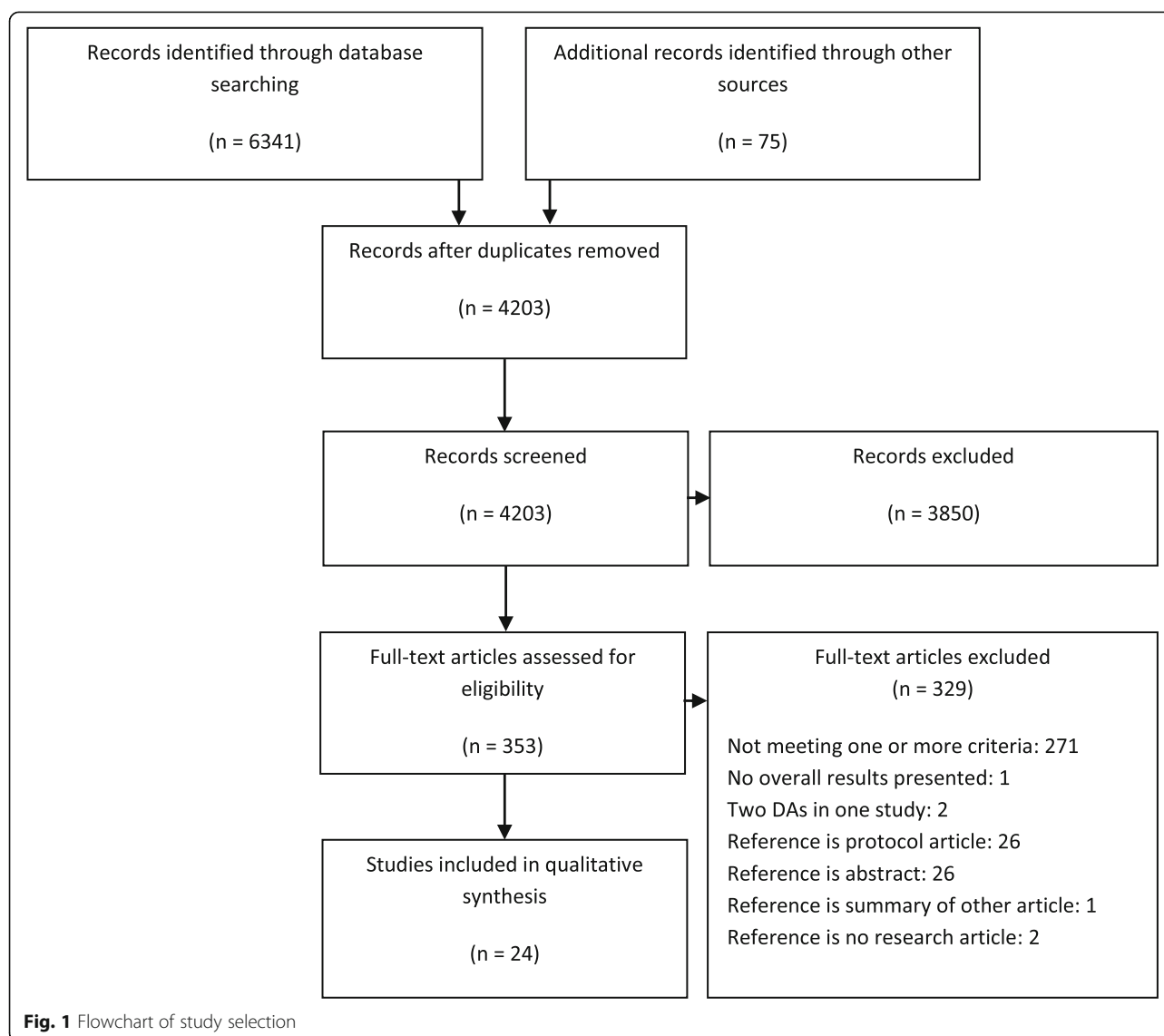


Table 2 SDM elements included in DAs and DA effects on SDM outcomes

DA study, year	SDM elements in DA					SDM outcomes				
	Situation diagnosis	Choice awareness	Option clarification	Harms and benefits	Patient preferences	Making decision	Knowledge	Patient participation	Decisional conflict	Satisfaction Time
Cardiovascular diseases										
Knops et al. 2014 [32]	✓	✓	✓	✓	✓	✓	↔	.	↔	.
Man-Son-Hing et al. 1999 [33]	?	?	✓	✓	✓	✓	.	↔	↔	.
Fraenkel et al. 2012 [34]	✓	?	✓	✓	✓
Thomas et al. 2013 [35]	✓	?	✓	✓	?	?	↔	.	↔	.
El-Jawahri et al. 2016 [36]	✓	✓	✓	?	?	?	↑	.	.	.
Kortland et al. 2017 [37]	✓	✓	✓	✓	✓	✓	.	.	↔	.
Thomson et al. 2007 [38]	?	?	✓	✓	✓	✓	↔	.	↑	.
Morgan et al. 2000 [39]	✓	?	✓	✓	?	?	↑	.	.	.
Coylewright et al. 2016 [40]	.	?	✓	✓	✓	✓	.	↔	↔	.
McAlister et al. 2005 [41]	✓	✓	✓	✓	✓	✓	.	.	↑	.
Respiratory diseases										
Gagné et al. 2017 [29]	✓	?	✓	✓	✓	✓	↔	.	↔	.
Slok et al. 2016 [42]	✓	?	✓	?	✓	✓
Diabetes										
Huang et al. 2017 [43]	✓	✓	✓	✓	✓	.	.	.	↔	.
Statin Choice [30, 31, 44, 45]	✓	?	✓	✓	.	✓	↑	↑	↑	↔
Mathers et al. 2012 [46]	?	?	✓	✓	✓	?	.	.	↑	↔
Heisler et al. 2014 [47]	✓	?	✓	✓	✓	✓	↔	.	↔	.
Bailey et al. 2016 [48]	✓	?	✓	✓	✓	✓	↑	.	↑	.
Denig et al. 2014 [49]	✓	✓	✓	✓	✓	✓
Diabetes Medication Choice [50, 51]	.	?	✓	✓	✓	.	↔	↑	↔	.
den Ouden et al. 2017 [52]	.	?	✓	✓	✓	✓

Elements: • = not present; ? = unclear; ✓ = present; Outcomes: • = not reported; ↔ = no statistically significant effect; ↑ = favored DA

and Additional file 5 their impact on SDM and health outcomes. We imputed the SD for the decisional conflict outcome for Mann et al. [30] using the SD found by Weymiller et al. [31] for the same outcome in the same context.

Discussion

This review presents an overview of chronic care DAs developed and tested in RCTs, SDM elements they support, and their effects on SDM outcomes and health outcomes. Most DAs support the clarification of options and the discussion of their benefits and harms, while other elements are less prevalent. Almost all trials were at an unclear or high risk of bias, and no association between SDM elements supported in the DA on the one hand and SDM outcomes achieved versus control on the other hand could be determined.

SDM elements handled by DAs

Our analysis of SDM elements supported is consistent with previous literature stating that most DAs focus and are tested on providing information or discussing choices rather than on creating empathic conversations [53]. We could not, however, estimate the relationship between the extent to which DAs support SDM elements and SDM outcomes.

Possibly, some SDM elements may have been left out of DAs by design. This choice may depend on what features were thought most important by the developers (e.g., patient education, risk communication, preference elicitation, or patient empowerment). The importance of incorporation of SDM elements in DAs may be situation-dependent, but the way this works is unclear. Future research should clarify this situation-dependence and eventually inform possible reconsideration of the IPDAS minimum standards for DA qualification [17].

DA effects

The inability to find any empiric association between features present and SDM outcomes prevents us from using this evidence base to make recommendations about the content of DAs for use in patients with chronic conditions. Multiple factors potentially explain the varying effects, including the following: whether a patient decision aid or conversation aid is used [10], chronicity of conditions [2], design process [54, 55], context, target population [19], and degree of detail needed [19]. Future studies may assess the dependency of DA effects on these factors and their interactions with the SDM elements.

Difficulties faced

Some difficulties were faced when conducting this review. A major difficulty during the article selection was

the suboptimal reporting of DA characteristics. The aim of DAs is not always explicitly described and if described, it still may be questionable whether implementing SDM is implicitly aimed for as the concept of SDM itself is highly debatable [56]. Namely, a review found 31 separate concepts to explicate SDM [57]. Our ability to categorize whether SDM elements were present was limited by the fact that some DAs were not available and/or the description of the DA's content was not clear and detailed. The latter is in line with the literature [58, 59]. Even when DAs were available and/or content was clearly described, it may not always be clear-cut whether or not an element is handled. Therefore, data regarding the SDM elements is based on reviewers' judgments. Furthermore, it may sometimes be unclear whether or not a condition is chronic (e.g., aneurysms). These conditions were included in this review in order to be as comprehensive as possible, but the decisions to be made may not be reversible over time or only to a limited extent. These aspects may have resulted in the fair inter-rater agreement. Another difficulty was found in the large methodological heterogeneity across studies (e.g., measurement instruments, timing of outcome measurements, and presentation of results).

More guidance is needed on the reporting of SDM elements and DA aims, the measurement instruments to use in RCTs studying DA effects, as well as the timing of outcome measurements and the way results are presented in articles. Furthermore, the quality of RCTs studying DA effects can be improved. The new Standards for UNiversal reporting of patient Decision Aid Evaluation studies (SUNDAE) checklist seems to meet this need as it helps to ensure the high-quality reporting of DA evaluation studies, as well as its intelligibility and transparency [59].

Strengths and limitations

This review is the first to report on SDM elements included in DAs developed for chronic conditions, and its relations to a range of SDM outcomes. This review underscores the importance of methodological improvement of DA evaluation studies, which hopefully will be attained by the new SUNDAE checklist [59].

Our review has some limitations. Since we were interested in the efficacy of DAs, we have limited our search strategy to RCTs [60], which may have led to exclusion of (well designed and developed) DAs that have not been tested in trials. Finally, we limited our search strategy to the most prevalent cardiovascular diseases, chronic respiratory diseases, and diabetes [22–24], an incomplete list of chronic diseases. This while a silver bullet of the literature probably brings to light what is happening in other chronic illnesses.

Future research

Future research should focus on empirically testing which SDM elements should be included in DAs, and take situation-dependency into account. This warrants studies with a sound methodology and low risk of bias that are currently lacking.

Conclusions

Tools to promote SDM for patients with chronic conditions support only some key recommended SDM elements thought to be important for SDM. The literature has not examined the relationship between explicit support for these elements in DAs and SDM outcomes.

Additional files

Additional file 1: PRISMA checklist. (DOC 63 kb)

Additional file 2: Search histories bibliographic databases. (DOCX 25 kb)

Additional file 3: Risk of bias assessment. (DOCX 46 kb)

Additional file 4: DA information. (DOCX 77 kb)

Additional file 5: DA effects. (DOCX 64 kb)

Abbreviations

95% CI: Confidence interval; DA: Decision aid; IPDAS: International Patient Decision Aid Standards; OR: Odds ratios; PRISMA: Preferred Reporting Items for Systematic Reviews and Meta-Analyses; PRO: Patient-reported outcomes; RCT: Randomized controlled trials; SD: Standard deviation; SDM: Shared decision making; SE: Standard error; SMD: Standardized mean difference; SUNDAE: Standards for Universal reporting of patient Decision Aid Evaluation studies

Acknowledgements

We would like to thank the Amsterdam Public Health research institute for providing a travel grant which enabled THW to visit the Mayo Clinic in Rochester (MN, USA). This grant provided the opportunity to work face-to-face on this review. We would like to thank Laxsini Murugesu and Anne Vloemans for their part in the abstract selection, as well as Miranda Romkes and Debora Mars-de Haan for their part in the data collection. Furthermore, we would like to thank Hans van der Wouden for his advice regarding methodological and practical issues.

Funding

Not applicable.

Availability of data and materials

The datasets used and analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

LJS and THW created the search strategy. LJS carried out the literature search. RR-G and THW selected abstracts and full-text articles and collected the data for selected articles resulting from the initial search strategy. MFS-H and THW selected abstracts and full-text articles resulting from the additional search strategy. GS-B, MFS-H, NRE, YZ-I, and THW collected the data for the selected full-text articles resulting from the additional search. GS-B decided upon conflicts in the selection of full-text articles, both resulting from the initial search and the additional search. OJP and THW assessed the risk of bias within studies on outcome level. RR-G, GS-B, MdW, OJP, MFS-H, NRE, YZ-I, MK, LJS, VMM, and FJS made substantial contributions to the manuscript by providing feedback on draft versions. THW processed this feedback, which resulted in the final version. All authors read and approved the final manuscript.

Ethics approval and consent to participate

Not applicable.

Consent for publication

Not applicable.

Competing interests

The Mayo Clinic Knowledge and Evaluation Research Unit produces and tests shared decision making interventions for patients with chronic conditions. Tools are made freely available at <http://shareddecisions.mayoclinic.org> and no income is generated from their distribution or use.

Publisher's Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Author details

¹Department of Medical Psychology, Amsterdam Public Health Research Institute, Amsterdam UMC, Vrije Universiteit Amsterdam, De Boelelaan 1117, Amsterdam, the Netherlands. ²Knowledge and Evaluation Research Unit, Mayo Clinic, Rochester, MN, USA. ³Division of Endocrinology, Department of Internal Medicine, "Dr. Jose E. González" University Hospital, Autonomous University of Nuevo Leon, Monterrey, Nuevo Leon, Mexico. ⁴Plataforma INVEST Medicina UANL-KER Unit Mayo Clinic, KER Unit México, "Dr. Jose E. González" University Hospital, Autonomous University of Nuevo Leon, Monterrey, Nuevo Leon, Mexico. ⁵Department of Medicine, Stanford University School of Medicine, Stanford, CA, USA. ⁶College of Public Health, Temple University, Philadelphia, PA, USA. ⁷Medical Decision Making, Department of Biomedical Data Sciences, Leiden University Medical Center, Leiden, the Netherlands. ⁸Medical Library, VU University, Amsterdam, the Netherlands.

Received: 22 November 2018 Accepted: 29 April 2019

Published online: 20 May 2019

References

- Hargraves I, LeBlanc A, Shah ND, Montori VM. Shared decision making: the need for patient-clinician conversation, not just information. *Health Aff.* 2016;35(4):627–9.
- Montori VM, Gafni A, Charles C. A shared treatment decision-making approach between patients with chronic conditions and their clinicians: the case of diabetes. *Health Expect.* 2006;9(1):25–36.
- Kunnenman M, Engelhardt EG, Ten Hove FL, Marijnen CA, Portielje JE, Smets EM, et al. Deciding about (neo-) adjuvant rectal and breast cancer treatment: missed opportunities for shared decision making. *Acta Oncol.* 2016;55(2):134–9.
- Stiggelbout AM, Pieterse AH, De Haes JCJM. Shared decision making: concepts, evidence, and practice. *Patient Educ Couns.* 2015;98(10):1172–9.
- Stiggelbout AM, Van der Weijden T, De Wit MP, Frosch D, Légaré F, Montori VM, et al. Shared decision making: really putting patients at the centre of healthcare. *BMJ.* 2012;344:e256.
- Barry MJ, Edgman-Levitan S. Shared decision making - the pinnacle of patient-centered care. *N Engl J Med.* 2012;366(9):780–1.
- Sullivan MD, Leigh J, Gaster B. Brief report: training internists in shared decision making about chronic opioid treatment for noncancer pain. *J Gen Intern Med.* 2006;21(4):360–2.
- Elwyn G, Edwards A, Hood K, Robling M, Atwell C, Russell I, et al. Achieving involvement: process outcomes from a cluster randomized trial of shared decision making skill development and use of risk communication aids in general practice. *Fam Pract.* 2004;21(4):337–46.
- IPDAS Collaboration. What are patient decision aids? (2017). <http://ipdas.ohri.ca/what.html>. Accessed 30 Oct 2018.
- Montori VM, Kunnenman M, Brito JP. Shared decision making and improving health care: the answer is not in. *JAMA.* 2017;318(7):617–8.
- Stacey D, Légaré F, Lewis K, Barry MJ, Bennett CL, Eden KB, et al. Decision aids for people facing health treatment or screening decisions. *Cochrane Database Syst Rev.* 2017;(4):CD001431.
- Durand MA, Carpenter L, Dolan H, Bravo P, Mann M, Bunn F, et al. Do interventions designed to support shared decision-making reduce health inequalities? A systematic review and meta-analysis. *PLoS One.* 2014;9(4):e94670.

13. Légaré F, Turcotte S, Stacey D, Ratté S, Kryworuchko J, Graham ID. Patients' perceptions of sharing in decisions. *Patient Patient Cent Outcomes Res*. 2012;5(1):1–19.
14. Dwamena F, Holmes-Rovner M, Gaulden CM, Jorgenson S, Sadigh G, Sikorski A, et al. Interventions for providers to promote a patient-centred approach in clinical consultations. *Cochrane Libr*. 2012;12:CD003267.
15. Joosten EA, DeFuentes-Merillas L, De Weert GH, Sensky T, Van Der Staak CPF, de Jong CA. Systematic review of the effects of shared decision-making on patient satisfaction, treatment adherence and health status. *Psychother Psychosom*. 2008;77(4):219–26.
16. Volk RJ, Llewellyn-Thomas H, Stacey D, Elwyn G. Ten years of the international patient decision aid standards collaboration: evolution of the core dimensions for assessing the quality of patient decision aids. *BMC Med Inform Decis Mak*. 2013;13(Suppl 2):S1.
17. Joseph-Williams N, Newcombe R, Politi M, Durand M-A, Sivell S, Stacey D, et al. Toward minimum standards for certifying patient decision aids: a modified Delphi consensus process. *Med Decis Mak*. 2014;34(6):699–710.
18. Kunnean M, Branda ME, Hargraves I, Pieterse AH, Montori VM. Fostering choice awareness for shared decision making: a secondary analysis of video-recorded clinical encounters. *Mayo Clin Proc*. 2018;2(1):60–8.
19. Stacey D, Légaré F, Col NF, Bennett CL, Barry MJ, Eden KB, et al. Decision aids for people facing health treatment or screening decisions. *Cochrane Database Syst Rev*. 2014;(1):CD001431.
20. Wieringa TH, Kunnean M, Rodriguez-Gutierrez R, Montori VM, de Wit M, Smets EM, et al. A systematic review of decision aids that facilitate elements of shared decision-making in chronic illnesses: a review protocol. *Syst Rev*. 2017;6(1):155.
21. Moher D, Liberati A, Tetzlaff J, Altman DG. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *Int J Surg*. 2010;8(5):336–41.
22. World Health Organization. Chronic obstructive pulmonary disease (COPD). 2017. <http://www.who.int/mediacentre/factsheets/fs315/en/>. Accessed 30 Oct 2018.
23. World Health Organization. Cardiovascular diseases (CVDs). 2017. <http://www.who.int/mediacentre/factsheets/fs317/en/>. Accessed 30 Oct 2018.
24. World Health Organization. Noncommunicable diseases. (2018). <http://www.who.int/mediacentre/factsheets/fs355/en/>. Accessed 30 Oct 2018.
25. Higgins JP, Altman DG, Gøtzsche PC, Jüni P, Moher D, Oxman AD, et al. The Cochrane Collaboration's tool for assessing risk of bias in randomised trials. *BMJ*. 2011;343:d5928.
26. Higgins JPT, Altman DG, Sterne JAC. Assessing risk of bias in included studies. In: Higgins JPT, Green S, editors. *Cochrane handbook for systematic reviews of interventions version 5.1.0* (updated March 2011). The Cochrane Collaboration; 2011.
27. The Cochrane Collaboration. Review Manager (RevMan) [Computer program]. Version 5.3. Copenhagen: The Nordic Cochrane Center; 2014.
28. Landis JR, Koch GG. The measurement of observer agreement for categorical data. *Biometrics*. 1977;33(1):159–74.
29. Gagné ME, Légaré F, Moisan J, Boulet L-P. Impact of adding a decision aid to patient education in adults with asthma: a randomized clinical trial. *PLoS One*. 2017;12(1):e0170055.
30. Mann DM, Ponienan D, Montori VM, Arciniega J, McGinn T. The statin choice decision aid in primary care: a randomized trial. *Patient Educ Couns*. 2010;80(1):138–40.
31. Weymiller AJ, Montori VM, Jones LA, Gafni A, Guyatt GH, Bryant SC, et al. Helping patients with type 2 diabetes mellitus make treatment decisions: statin choice randomized trial. *Arch Intern Med*. 2007;167(10):1076–82.
32. Knops A, Goossens A, Ubbink D, Balm R, Koelemay M, Vahl A, et al. A decision aid regarding treatment options for patients with an asymptomatic abdominal aortic aneurysm: a randomised clinical trial. *Eur J Vasc Endovasc Surg*. 2014;48(3):276–83.
33. Man-Son-Hing M, Laupacis A, O'Connor AM, Biggs J, Drake E, Yetisir E, et al. A patient decision aid regarding antithrombotic therapy for stroke prevention in atrial fibrillation: a randomized controlled trial. *JAMA*. 1999;282(8):737–43.
34. Fraenkel L, Street RL, Towle V, O'leary JR, Iannone L, Ness PH, et al. A pilot randomized controlled trial of a decision support tool to improve the quality of communication and decision-making in individuals with atrial fibrillation. *J Am Geriatr Soc*. 2012;60(8):1434–41.
35. Thomas KL, Zimmer LO, Dai D, Al-Khatib SM, LaPointe NMA, Peterson ED. Educational videos to reduce racial disparities in ICD therapy via innovative designs (VIVID): a randomized clinical trial. *Am Heart J*. 2013;166(1):157–63.
36. El-Jawahri A, Paasche-Orlow MK, Matlock D, Stevenson LW, Lewis EF, Stewart G, et al. Randomized, controlled trial of an advance care planning video decision support tool for patients with advanced heart failure. *Circulation*. 2016;134(1):52–60.
37. Kortland NM, Ahmed Y, Koolbergen DR, Brouwer M, de Heer F, Klijn J, et al. Does the use of a decision aid improve decision making in prosthetic heart valve selection?: a multicenter randomized trial. *Circ Cardiovasc Qual Outcomes*. 2017;10(2):e003178.
38. Thomson RG, Eccles MP, Steen IN, Greenaway J, Stobart L, Murtagh MJ, et al. A patient decision aid to support shared decision-making on anti-thrombotic treatment of patients with atrial fibrillation: randomised controlled trial. *BMJ Qual Saf*. 2007;16(3):216–23.
39. Morgan MW, Deber RB, Llewellyn-Thomas HA, Gladstone P, Cusimano R, O'Rourke K, et al. Randomized, controlled trial of an interactive videodisc decision aid for patients with ischemic heart disease. *J Gen Intern Med*. 2000;15(10):685–93.
40. Coylewright M, Dick S, Zmolek B, Askelin J, Hawkins E, Branda M, et al. PCI choice decision aid for stable coronary artery disease: a randomized trial. *Circ Cardiovasc Qual Outcomes*. 2016;9(6):767–76.
41. McAlister FA, Man-Son-Hing M, Straus SE, Ghali WA, Anderson D, Majumdar SR, et al. Impact of a patient decision aid on care among patients with nonvalvular atrial fibrillation: a cluster randomized trial. *Can Med Assoc J*. 2005;173(5):496–501.
42. Slok AH, Kotz D, van Breukelen G, Chavannes NH, Rutten-van Mölken MP, Kerstjens HA, et al. Effectiveness of the assessment of burden of COPD (ABC) tool on health-related quality of life in patients with COPD: a cluster randomised controlled trial in primary and hospital care. *BMJ Open*. 2016; 6(7):e011519.
43. Huang ES, Nathan AG, Cooper JM, Lee SM, Shin N, John PM, et al. Impact and feasibility of personalized decision support for older patients with diabetes: a pilot randomized trial. *Med Decis Mak*. 2017;37(5):611–7.
44. Nannenga MR, Montori VM, Weymiller AJ, Smith SA, Christianson TJ, Bryant SC, et al. A treatment decision aid may increase patient trust in the diabetes specialist. The statin choice randomized trial. *Health Expect*. 2009;12(1):38–44.
45. Perestelo-Pérez L, Rivero-Santana A, Boronat M, Sánchez-Afonso JA, Pérez-Ramos J, Montori VM, et al. Effect of the statin choice encounter decision aid in Spanish patients with type 2 diabetes: a randomized trial. *Patient Educ Couns*. 2016;99(2):295–9.
46. Mathers N, Ng CJ, Campbell MJ, Colwell B, Brown I, Bradley A. Clinical effectiveness of a patient decision aid to improve decision quality and glycaemic control in people with diabetes making treatment choices: a cluster randomised controlled trial (PANDAS) in general practice. *BMJ Open*. 2012;2(6):e001469.
47. Heisler M, Choi H, Palmisano G, Mase R, Richardson C, Fagerlin A, et al. Comparison of community health worker-led diabetes medication decision-making support for low-income Latino and African American adults with diabetes using e-health tools versus print materials: a randomized, controlled trial. *Ann Intern Med*. 2014;161(10_Supplement):S13–22.
48. Bailey RA, Pfeifer M, Shillington AC, Harshaw Q, Funnell MM, VanWingen J, et al. Effect of a patient decision aid (PDA) for type 2 diabetes on knowledge, decisional self-efficacy, and decisional conflict. *BMC Health Serv Res*. 2016;16(1):10.
49. Denig P, Schuling J, Haaijer-Ruskamp F, Voorham J. Effects of a patient oriented decision aid for prioritising treatment goals in diabetes: pragmatic randomised controlled trial. *BMJ*. 2014;349:g5651.
50. Mullan RJ, Montori VM, Shah ND, Christianson TJ, Bryant SC, Guyatt GH, et al. The diabetes mellitus medication choice decision aid: a randomized trial. *Arch Intern Med*. 2009;169(17):1560–8.
51. Karagiannis T, Liakos A, Branda ME, Athanasiadou E, Mainou M, Boura P, et al. Use of the diabetes medication choice decision aid in patients with type 2 diabetes in Greece: a cluster randomised trial. *BMJ Open*. 2016;6(11): e012185.
52. Den Ouden H, Vos RC, Rutten GE. Effectiveness of shared goal setting and decision making to achieve treatment targets in type 2 diabetes patients: a cluster-randomized trial (OPTIMAL). *Health Expect*. 2017;20(5):1172–80.
53. Serrano V, Rodriguez-Gutierrez R, Hargraves I, Gionfriddo M, Tamhane S, Montori V. Shared decision-making in the care of individuals with diabetes. *Diabet Med*. 2016;33(6):742–51.
54. Feldman-Stewart D, Brennenstuhl S, Brundage MD. A purpose-based evaluation of information for patients: an approach to measuring effectiveness. *Patient Educ Couns*. 2007;65(3):311–9.

55. Witteman HO, Dansokho SC, Colquhoun H, Coulter A, Dugas M, Fagerlin A, et al. User-centered design and the development of patient decision aids: protocol for a systematic review. *Syst Rev*. 2015;4(1):11.
56. Krones T, Keller H, Becker A, Sönnichsen A, Baum E, Donner-Banzhoff N. The theory of planned behaviour in a randomized trial of a decision aid on cardiovascular risk prevention. *Patient Educ Couns*. 2010;78(2):169–76.
57. Makoul G, Clayman ML. An integrative model of shared decision making in medical encounters. *Patient Educ Couns*. 2006;60(3):301–12.
58. Lewis KB, Wood B, Sepucha KR, Thomson RG, Stacey D. Quality of reporting of patient decision aids in recent randomized controlled trials: a descriptive synthesis and comparative analysis. *Patient Educ Couns*. 2017;100(7):1387–93.
59. Sepucha KR, Abhyankar P, Hoffman AS, Bekker HL, LeBlanc A, Levin CA, et al. Standards for UNiversal reporting of patient decision aid evaluation studies: the development of SUNDAE checklist. *BMJ Qual Saf*. 2017;0:1–9.
60. Bouter LM, van Dongen MCJM, Zielhuis GA. *Epidemiologisch onderzoek: Opzet en interpretatie*. 6th ed. Houten: Bohn Stafleu Van Loghum; 2010.

Ready to submit your research? Choose BMC and benefit from:

- fast, convenient online submission
- thorough peer review by experienced researchers in your field
- rapid publication on acceptance
- support for research data, including large and complex data types
- gold Open Access which fosters wider collaboration and increased citations
- maximum visibility for your research: over 100M website views per year

At BMC, research is always in progress.

Learn more biomedcentral.com/submissions

